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PARVERY-SUIVAX Vaccino inattivato in emulsione iniettabile per suini

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Porcine parvovirus, Inactivated

Product identification

Medicine name:
PARVERY-SUIVAX Vaccino inattivato in emulsione iniettabile per suini

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Porcine parvovirus, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, serotype 2, Inactivated
50.00 international unit(s) / 2.00 millilitre(s)

Porcine parvovirus, Inactivated

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
28/04/2004

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
3/11/2009

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet